

TITLE**POLYAMIDE AND POLYVINYLBUTYRAL COMPOSITIONS AND
BLENDS HAVING ENHANCED SURFACE PROPERTIES AND
ARTICLES MADE THEREFROM**

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This application claims the benefit of U.S.
Provisional Application No. 60/454,890, filed March 14,
2003.

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FIELD OF THE INVENTION

The present invention relates to polyamide blends
with polyvinylbutyral (PVB). More particularly, the
present invention relates to such blends, processes for
the manufacture of such materials, and molded articles
15 prepared therefrom.

BACKGROUND OF THE INVENTION

It is well known that toughening agents such as
grafted rubbers can be employed to improve the
20 toughness of polyamides. See generally, U.S. Pat. No.
4,174,358. It is also well known that glass fibers can
be incorporated into such polyamide blends to increase
their stiffness. Such a product has been available
from E.I. DuPont de Nemours and Company (DuPont) for a
25 number of years, under the name ZYTEL® 80G33HS1L BK104.
It is also well known that plasticized polyvinylbutyral
(PVB) can be used as a toughener in, for example, 6-
nylon. See generally, U.S. Pat. No. 5,770,654 which is
directed to such compositions suitable for a variety of
30 applications where good toughness is required, as in
packaging subjected to rough handling. As used herein
polyvinylbutyral is abbreviated as "PVB".

Plasticized PVB can be difficult to handle as a feed to a compounding extruder due to its inherent stickiness. Similarly PVB sheet is a material that can be difficult to work with because of the tendency to
5 adhere to itself. Sheets of PVB can stick together, or bind, with such strength that it is very difficult to separate the layers. The irreversible nature of this self-adhesion by PVB is referred to in the art of PVB manufacture as "blocking". Once PVB "blocks", process
10 difficulties are encountered. This tendency to block can make manufacturing processes that incorporate PVB complex and difficult. Consequently, continuous processes in which PVB is handled either in sheet form or in small shredded pieces can be very expensive to
15 run, and therefore are not practical.

Moreover, blends of PVB sheet or small shredded pieces with other materials can block in the same manner as homogenous PVB compositions. Such blends of PVB with other polymers can be difficult to obtain in a
20 cost-effective manner. A preferred process for preparing blends of PVB with other polymers would utilize conventional loss-in-weight screw feeders, which are found throughout that industry.

Recent work in the field indicates that blends of
25 PVB with polyethylene and grafted rubbers are sufficiently non-sticky that they can be fed into a compounding extruder. See for example, WO 02/12356 directed to a process for preparing pellets from PVB scrap material.

30 It is an object of the present invention to provide PVB-toughened polyamide compositions having

enhanced surface adhesion, making them suitable for use in a variety of applications.

SUMMARY OF THE INVENTION

In one aspect, the present invention is a
5 polyamide composition having enhanced surface adhesion properties, comprising: the reaction product of a silane coupling compound and a polyamide blend.

In another aspect, the present invention is a thermoplastic polyamide composition comprising: (a)
10 from about 5 to about 30 weight percent of a free-flowing toughener comprising from about 20 weight percent to about 95 weight percent polyvinyl butyral; (b) complementally, 95 to 25 weight percent polyamide that is melt processible below about 320°C and a number
15 average molecular weight of at least 5,000; (c) a coupling agent; and (d) optionally, a filler in an amount of up to about 45 weight percent.

In still another aspect, the present invention is a process for enhancing the surface adhesion of a
20 polyamide blend comprising the step: coating a surface of the polyamide blend with a silane coupling agent.

DETAILED DESCRIPTION OF THE INVENTION

In one embodiment, the present invention is a
25 thermoplastic polyamide composition having enhanced surface adhesion properties. Enhanced surface adhesion in a polyamide of the present invention is determined relative to conventional polyamide compositions. A polyamide composition of the present invention is
30 preferably a blend of a polyamide with at least one other polymer. The polyamide composition of the

present invention can comprise, in addition to the polyamide, an adhesion-promoting polymer. In one embodiment the polyamide is preferably a blended composition of a polyamide with at least one
5 thermoplastic polymer such as is described in U.S. Pat. No. 4,174,358. Such blends are commercially available from DuPont under the trade name Zytel®, for example, or alternatively a polyamide/PVB blend such as is described hereinbelow.

10 Polyamide blends suitable for use in the practice of the present invention are blends of conventional polyamides such as: Nylon 6; Nylon 66; Nylon 69; Nylon 610; and Nylon 612; Nylon 11; Nylon 12; Nylon 12, 12; and copolymers of epsilon-caprolactam with
15 hexamethylenediamine and adipic acid for example, with at least one polymer of the type described in U.S. Pat. No. 4,174,358. For example, suitable polyamides, as described in U.S. Pat. No. 4,174,358, can be blended with thermoplastic or elastomeric polymers as
20 tougheners such as polyolefins, ethylene copolymers, grafted polymers and copolymers such as maleic anhydride grafted ethylene copolymers, and similar polymers or mixtures thereof. A more complete list of suitable elastomeric or thermoplastic polymers can be
25 found in the above-referenced U.S. patent. The blends can comprise from about 1 to about 99 wt% of the polyamide, preferably from about 60 to about 99 wt%, and more preferably from about 80 to about 95 wt% polyamide, by weight of the total composition.

30 The polyamide blend can comprise on its surface, or incorporated within the polymer matrix, an adhesion-

promoting compound. Preferably, the adhesion promoting compound is a silane coupling compound, as described further hereinbelow.

In a preferred embodiment, the present invention
5 is a toughened polyamide composition having enhanced surface adhesion properties with or without use of a coupling compound. A composition of the present invention comprises a free-flowing PVB composition as a toughener, as described in WO 0212356, incorporated
10 herein by reference. A composition of the present invention comprises from about 5 wt% to about 30 wt%, preferably from about 5 wt% to about 28 wt%, more preferably from about 6 wt% to about 25 wt%, and most preferably from about 7 wt% to about 25 wt% of a free-
15 flowing PVB composition. The toughener comprises from about 20 to about 95 wt%, preferably from about 40 wt% to about 95 wt%, more preferably from about 60 wt% to about 95 wt%, and most preferably from about 75 wt% to about 95 wt% PVB. The compositions and blends of this
20 invention can be prepared by mixing or blending a free-flowing toughener with nylon, an optional coupling agent, and other ingredients to produce a toughened polyamide blend having enhanced surface properties.

The toughener comprises at least one component in
25 addition to the PVB. Such other components can be monomeric or polymeric materials, or mixtures thereof. The other components can be selected from polymers and/or monomers that have reactive functionality, or non-reactive polymer and/or monomers such as, for
30 example, polyethylene, polypropylene, polyvinylchloride, nylon, other thermoplastic

materials, or mixtures thereof. Preferably the second component is a polymer composition that includes reactive functionality such as anhydride functionality, such as is available commercially from E. I. DuPont de Nemours and Company under the Fusabond® brand name, or carboxylic acid functionality. Fusabond® polymers are polyolefins having anhydride functionality. The other components are present in the toughener in amounts that are complimentary to the amount of PVB in the toughener, that is to bring the total percentage of PVB and other component(s) to 100 wt%.

The polyamide with which the toughener is blended can be any amorphous or crystalline polyamide as described in U.S. Pat. No. 5,770, 654, for example, or in U.S. Pat. No. 4,174,358. Preferably, the polyamide is melt processible below a temperature of about 320°C and has a number average molecular weight of at least 5,000. The polyamide component can be present in an amount of from about 25 wt% to about 95 wt%. Preferably, the polyamide component is present in an amount of from about 30 wt% to about 90 wt%, more preferably from about 40 wt% to about 90 wt%, most preferably from about 50 wt% to about 90 wt%.

Fillers can optionally be present in an amount of from about 0 to about 45 wt%. If included, fillers are preferably present in an amount of from about 1 wt% to about 45 wt%. Suitable fillers are, for example, calcined clay, metal carbonates, titanium dioxide, wollastonite, glass, or talc. Glass-filled compositions can include glass from any number of sources, or in any form. For example, glass can be

included in crushed form, as shards, as particulate glass, as fiber, or any other form that glass can be included and processed using the process described herein.

5 As noted hereinabove a coupling agent can optionally be used in the composition of the present invention. The coupling agent can further enhance the adhesive surface properties of the toughened polyamide compositions of the present invention. The coupling
10 agent can be a silane compound. Preferably the coupling compound is selected from the group consisting of: gamma-aminopropyltrimethoxysilane; gamma-aminopropyltriethoxysilane; N-2-aminopropyltrialkoxysilane; or N-(2-aminoethyl)-3-
15 aminopropylmethyldialkoxysilane. The coupling compound can be present in an amount of at least about 0.01 wt%. Preferably, the coupling agent is present in an amount of from about 0.1 to about 3 wt%. More preferably, the coupling agent is present in an amount of from about
20 0.3 wt% to about 2.0 wt%, and most preferably in an amount of from about 0.5 wt% to about 1.5 wt%.

 An antioxidant is not required, however an antioxidant can be preferred. If included, the antioxidant can be present in an amount of at least
25 about 0.1% by weight, and up to an amount where the effect of the antioxidant is optimal.

 In another embodiment, the present invention is a process for preparing polyamide compositions of the present invention having enhanced adhesion. The
30 polyamide of the present invention can be obtained by

blending a suitable polyamide with PVB, for example Ecocite™.

Alternatively, polyamides having enhanced adhesive properties are obtained by further incorporating a
5 coupling or crosslinking agent with the toughened polyamide or a suitable polyamide blend. For example, a silane coupling agent can be incorporated by either inclusion into the bulk of the polyamide composition, or by coating the surface of the polyamide composition.
10 The coupling compound can be incorporated in either manner as an aqueous solution. The pH of the solution can be lowered using an acid such as acetic acid or citric acid, for example.

In a preferred embodiment, the present invention
15 is a process wherein plasticized PVB and three other ingredients (a reactive polymer such as Fusabond®, a non-reactive polymer such as polyethylene, polypropylene, or ethylene/n-butyl, and an antioxidant) are made into pellets and subsequently combined with a
20 polyamide to provide the toughened polyamide compositions of the present invention.

The toughener can be prepared prior to addition to the polyamide, for example, by mixing the components at an elevated temperature in the range of from about
25 100°C to about 280°C, preferably from about 150°C to about 220°C to provide a homogeneous melt blend. The blend obtained from the mixing procedure can be transferred by some means to a set of roll mills for additional mixing and to press the blend into a sheet
30 form. A strip of the sheet can be fed either continuously or by a batch process to an extruder, but

preferably the sheet is continuously fed using, for example, a belt feeder. Once inside the extruder, the sheet is melted and the melt is filtered to remove solid contamination. The polymer can be pelletized by
5 any known or conventional method. For example, the filtered melt can be distributed to a die wherein the die has multiple holes. In such a process the melt exits the die at the die face, which can be positioned just above the surface of the water in a tank filled
10 with water, or submerged under the surface of the water to quickly cool (quench) the melt as it exits the die. An under water face cutter can be used to cut the polymer exiting the die face to form pellets. The water quenches the pellets and carries them to a filter
15 screen to separate them from the bulk water. The wet pellets can be dried, for example in a fluidized dryer, before they are packed.

The pellets thus obtained can be mixed by melt blending with suitable polyamide compositions, as
20 described in U.S. Pat. No 5,770,654, herein incorporated by reference. For example, the toughened polyamide blends of the present invention can be obtained by melt blending, or melt mixing in any suitable blending or mixing device, such as a Banbury
25 blenders, Haake mixers, Farrell mixers, or extruders. Extruders can be either single screw or twin screw extruders with screws having various degrees of severity. Mixing or blending can be done at a temperature in the range of from about 200°C to about
30 320°C, and preferably at a temperature in the range of from about 230°C to about 300°C. The blends can be palletized by any known conventional method. Preferably

pellets are formed by cutting extruded strands of the blend.

Still alternatively, a suitable polyamide composition can be obtained by a process comprising the
5 step of blending a free-flowing PVB composition, such as Ecocite™ with a polyamide.

In another embodiment, the present invention is an article obtained from the polyamide compositions of the present invention. Articles of the present invention
10 include laminate articles, shaped articles, etc. Laminates comprising the polyamide compositions of the present invention can be incorporated into various other articles such as, for example, cars, trains, automobiles, appliances, boats, acoustic tiles,
15 acoustic flooring, walls, ceilings, roofing materials or other articles where sound damping and/or tough polymers are desirable.

Unlike conventional PVB products, the polyamide compositions of the present invention are not fully
20 transparent and therefore are not suitable for use in glazing, in automobile windshields, or similar applications where transparency is an important factor. However, the toughened polyamide compositions of the present invention have enhanced impact performance, as
25 indicated by Notched Izod tests. As such, the polyamide compositions of the present invention can be used to produce articles that can be used in applications requiring tough polymeric structural components.

30 The enhanced adhesive properties of the polyamides of the present invention make the polyamides of the

present invention suitable for use in laminate articles. In a particularly preferred embodiment, polyamide compositions of the present invention can be laminated to other polymeric materials, such as
5 thermoplastic elastomers (TPEs), conventional PVB, polyurethane, polyvinylchloride, polycarbonate, or other polyamides.

TPEs are thermoplastic materials that have rubber-like properties and are polymers which are soft to the
10 touch. However, TPEs do not generally have good adhesion to rigid polymers. TPE laminates with polyamides of the present invention can reduce or eliminate this adhesion problem in many cases. Such multilayer constructions of TPE's with the polyamides
15 of the present invention can find use in automobile interiors, buttons or switches on electronic equipment or electronic devices such as stereos, compact disc players, telephones, television sets, remote controls, computers, keypads, touch-screens, for example.

20 In another preferred embodiment, the polyamide compositions of the present invention can be laminated with PVB to yield PVB laminates having substantial sound reduction properties. Sound reducing panels can find use in many applications, including use in
25 construction materials, appliance panels, automobile panels, highway sound barriers, walls, ceilings, and floors, for example.

In a particularly preferred embodiment, a single sheet of PVB can be laminated between two polyamide
30 sheets of the present invention. Laminates having at least two sheets of the polyamide of the present

invention adhered on opposite surfaces of a PVB polymer sheet can demonstrate improved structural strength compared to one sheet of the polyamide having twice the thickness of one of the laminated polyamide sheets.

5 Such laminates can find use in car door panels, boat hulls, or other similar uses to impart structure, strength, and noise reduction. Lamination of the sheets can be by conventional methods, such as coextrusion, press molding, or injection molding. The
10 laminates can be hard panels or soft sheets for use in a variety of applications.

In still another embodiment the polyamide compositions of the present invention can be used as an adhesive layer or film for the purpose of binding glass
15 fibers that are on or near the surface of articles comprising fiber-glass filled polyamide compositions. Glass-filled polyamide compositions can have problems which result from loss of adhesion between the polyamide and the glass fibers on or near the surface
20 of the glass-filled polyamide composition. This can in turn lead to shedding of glass fibers and loss of same to the surrounding environment, or unwanted transfer of glass fibers to other articles or people. A glass-filled polyamide composition of the present invention
25 can demonstrate improved adhesion to glass fibers at or near the surface, and therefore can reduce the occurrences of problems associated with loss of adhesion between glass fibers and glass-filled polyamide compositions.

EXAMPLES

Examples E1 & E2 and Comparative Example C1

Extrusion Process to Produce Polymer Blends

5 ECOCITE™, commercially available from DuPont, was melt blended together with nylon-6 (Ultradid® B-3, available commercially from BASF Corp.). The comparative example C1 included no Ecocite™, but
10 instead included a mixture of conventional tougheners (T-mix) at 15 wt%. During the operation for melt blending the ingredients were charged to the blender using individually controlled loss in weight feeders. The mixture was compounded by melt blending in a 40mm
15 Werner & Pfleiderer co-rotating twin screw extruder with a barrel temperature about 240°C and a die temperature of about 260°C. All of the ingredients were fed into the first barrel section. Extrusion was carried out with a port under vacuum. The screw speed
20 was 250 rpm and the total extruder feed rate was 150 pounds per hour.

The resulting strand was quenched in water, cut into pellets, and sparged with nitrogen until cool.
25 The moisture in the resulting pellets was adjusted to between 0.1% and 0.2% by drying or adding additional water as required.

Modified Compressive Shear Stress (CSS) Test for Adhesion of Laminated Polymer Plate

30 Plagues of 5x5 inch with 2mm thickness were molded in an injection molding machine according to ISO 294. Stiff PVB (that is, PVB having less than 30 pph of plasticizer) was sandwiched between two plaques in a
35 humidity controlled room. After being autoclaved at 135°C for 20 minutes, the 5x5 inch laminated polymer plate was cut to obtain six 1x1 inch squares from the center plate. Each square was sheared at 45 degree
40 angle in an Instron in a humidity controlled room. Force (pounds per inch) required to shear the square apart (CSS) was measured and recorded. The average force (Avg) and the standard deviation was calculated for each sample and recorded in Table 1. The amount of
45 ECOCITE™ Grade H used in Examples 1 and 2 (E1 & E2) was varied. For comparative example 1 (C1), a mixture (T-mix) of conventional tougheners -- Fusabond® (available

from DuPont) and Engage® (available from DuPont-Dow Elastomers) was included.

Table 1

Example #	Additive (Wt%)	CSS Avg (Std. Dev)
C1	T-mix (15)	1345.318 (360.322)
E1	ECOCITE™ H (15)	1859.696 (156.49)
E2	ECOCITE™ H (25)	1736.568 (214.004)

5

Replacing the conventional tougheners with ECOCITE™ H resulted in increased adhesion in the laminated polymer plates.

10 Examples E3 to E11

The same process and procedures in above Examples E1, E2, & C1 were used for Examples E3 to E11 in Table 2 except that the Stiff PVB was coated with Silquest A-1100® using each of three coating methods before being dried and laminated:

Coating Method 1 - 3% Silquest® A-1100 aqueous solution at pH 7
 Coating Method 2 - 3% Silquest® A-1100 aqueous solution at pH 4.0 with Acetic Acid
 Coating Method 3 - 3% Silquest® A-1100 aqueous solution at pH 3.0 with Citric Acid

25

Table 2 Improved Adhesion on Nylon 6 Blends with Silane Coating			
Example #	Additive (Wt%)	Method	CSS Avg (Std. Dev)
E3	T-mix (15)	1	2210.537 (54.441)
E4	ECOCITE™ H (15)	1	1820.596 (578.155)
E5	ECOCITE™ H (25)	1	1799.268 (411.985)
E6	T-mix(15)	2	2161.352 (108.387)
E7	ECOCITE™ H (15)	2	2536.738 (77.387)
E8	ECOCITE™ H (25)	2	2344.607 (73.163)
E9	T-mix(15)	3	2221.338 (71.298)
E10	ECOCITE™ H (15)	3	2476.787 (132.555)
E11	ECOCITE™ H (25)	3	2303.907 (58.886)

The adhesion was increased by addition of Silquest® A-1100 in each of the three different ways.

Examples E12 to E15 & C2

The same process and procedures in above Examples E1, E2, & C1 were used for Examples E12 to E15 & C2 in Table 3 except 33 wt% glass fiber (PPG 3660, available commercially from PPG Industries) was fed into the sixth barrel section of the extruder by use of a side feeder. Adhesion results are reported in Table 3.

Table 3

Example #	Additive (Wt%)	CSS Avg (Std. Dev)
C2	T-mix(13)	871.371 (281.556)
E 12 ^a	ECOCITE™ E (13)	2005.93 (292.506)
E 13 ^a	ECOCITE™ G (13)	2664.57 (149.503)
E 14 ^a	ECOCITE™ H (13)	2637.45 (137.638)
E15 ^b	ECOCITE™ H (10.5)	2603.35 (179.75)

^anatural color glass-filled nylon 6

^bblack glass-filled nylon 6

Replacing conventional tougheners with ECOCITE™ Grade E, G, or H resulted in increased adhesion in the laminated polymer plates as E12 to E15 are compared to C2.

Examples E16 to E18 and C3

The same process and procedures in above Examples E12 to E14, & C2 were used for Examples E16 to E18 and C3 in Table 4 except that stiff PVB was replaced by "standard" Butacite® B140C, available from DuPont. Adhesion results are reported in Table 4.

Table 4

Example #	Additive (13 Wt%)	CSS Avg (Std. Dev)
C3	T-mix	764.1 (100.1)
E16	ECOCITE™ E	1130.3 (59.9)
E17	ECOCITE™ G	1140.6 (161.7)
E18	ECOCITE™ H	1120.3 (65.6)

Replacing conventional tougheners with ECOCITE™ Grade E, G, or H resulted in increased adhesion in the laminated polymer plates as E16 to E18 are compared to C3.

Examples E19 to E24

5 The same process and procedures in above Examples E18, & C3 were used for Examples E19 to E24 in Table 5 except standard Butacite® B140C was used in place of stiff PVB and coated with Silquest® A-1100 using each of three coating methods:

- 10 Coating Method 1 - 3% Silquest® A-1100 aqueous solution at pH 7
Coating Method 2 - 3% Silquest® A-1100 aqueous solution at pH 4.0 with Acetic Acid
Coating Method 3 - 3% Silquest® A-1100 aqueous solution at pH 3.0 with Citric Acid

The results are reported in Table 5.

Table 5 Improved Adhesion on Glass Filled Nylon 6 Blends with Silane Coating on Standard Butacite®			
Example #	Additive (13 Wt%)	Method	CSS Avg (Std. Dev)
E19	T-mix	1	916.472 (165.416)
E20	ECOCITE™ H	1	1282.85 (306.367)
E21	T-mix	2	1499.83 (104.3)
E22	ECOCITE™ H	2	1934.7 (305.721)
E23	T-mix	3	1580.22 (128.283)
E24	ECOCITE™ H	3	2227.3 (279.978)

- 5 The adhesion was increased by addition of Silane in each of the three different coating methods.

Examples E25, E26, C5 & C6

- 10 Using stiff PVB, the same process and procedures in above Examples E1, E2, & C1 were used for Examples E25, E26, & C6 in Table 6 except Nylon 6 was replaced by Zytel® 101 (available commercially from E.I. DuPont de Nemours and Company) and 40 wt% mineral (Translink
- 15 HF900, available commercially from Englehard) was fed into the sixth barrel section of the extruder by use of a side feeder. Temperatures for extruder barrel and die were increased to about 280°C and 290°C, respectively. Zytel® 11C40, commercially available
- 20 from DuPont, was used to make the laminated plate in Control example C5. Adhesion results are reported in Table 6.

Table 6 Effect of ECOCITE™ on Mineral Filled Nylon 66		
Example #	Additive (Wt%)	CSS Avg (Std. Dev)
C5	Zytel® 11C40	1445.4 (213.27)
C6	FA ^a (9)	1088.5 (226.48)
E25	ECOCITE™ H (18)	2095.1 (84.29)
E26	ECOCITE™ H (9) + FA (9)	2297.9 (240.1)

^aFusabond® A MG-423D

- 25 The adhesion was increased by addition of ECOCITE™ H.

Examples E27 to E38

The same process and procedures in above Examples E25, E26, C5, & C6 were used for Examples E27 to E38 in Table 7 except Stiff PVB was coated with Silane such as Silquest® A-1100 in three ways before being dried and laminated:

- Coating Method 1 - 3% Silquest® A-1100 aqueous solution at pH 7
- Coating Method 2 - 3% Silquest® A-1100 aqueous solution at pH 4.0 with Acetic Acid
- Coating Method 3 - 3% Silquest® A-1100 aqueous solution at pH 3.0 with Citric Acid
- The results are reported in Table 7.

Table 7 Improved Adhesion on Mineral Filled Nylon 6 Blends			
Example #	Additive (Wt%)	Method	CSS Avg (Std. ev)
E27 ^a	Zytel® 11C40	1	3334.7 (252.27)
E28	FA ^b (9)	1	2597.1 (52.81)
E29	ECOCITE™ H (18)	1	2480 (154.42)
E30	ECOCITE™ H (9)+FA (9)	1	2491.2 (90.1)
E31 ^a	Zytel® 11C40	2	2843.1 (148.18)
E32	FA (9)	2	2291.8 (186.42)
E33	ECOCITE™ H (18)	2	2327.2 (154.51)
E34	ECOCITE™ H (9)+FA (9)	2	2272.4 (216.95)
E35 ^a	Zytel® 11C40	3	2879.8 (124.33)
E36	FA (9)	3	2293.3 (109.65)
E37	ECOCITE™ H (18)	3	2074.5 (138.71)
E38	ECOCITE™ H (9)+FA (9)	3	2476.4 (135.71)

^aSample is 100% Zytel 11C40, commercially available from Dupont

^bFusabond® A MG-423D

Comparing Examples E27 to E38 versus Examples E25, E26, C5, & C6, respectively, the adhesion was increased by addition of Silquest® A-1100 by all methods.

Examples E39 to E44: Enhanced Adhesion Through Extruder Silane Addition for Mineral Filled Nylon 66

The same process and procedures in above Examples E25, E26, & C6 were used for Examples E39 to E44 with stiff PVB in Table 8 except Silane such as Silquest A-

1100® was mixed with the Zytel® 101 before feeding to the extruder. Adhesion results are reported in Table 8.

Table 8 Improved Adhesion on Mineral Filled Nylon 6 Blends with Silane Addition^a to Polymer in the Extruder		
Example #	Additive (Wt%)	CSS Avg (Std. Dev)
E39	ECOCITE™ H (9)+Si	3247.8 (77.384)
E40	ECOCITE™ H (12)+Si	3369.8 (260.443)
E41	ECOCITE™ H (18)+Si	3384.5 (298.798)
E42	ECOCITE™ H (9)+FA (9) +Si	2938.5 (92.315)
E43	FA ^b (9)+ Si	2820.7 (181.175)
E44	FC ^c (9) + Si	2760.9 (208.301)

5 ^aThe Silane compound is present at a level of less than 0.5 wt% in each sample.

^bFusabond® A MG-423D

^cFusabond® C MC190D (commercially from E.I. DuPont de Nemours and Company.

10

 The adhesion was increased by addition of Silquest® A-1100 in the extruder.

15 SEM (Scanning Electronic Microscopy) Pictures of Example E15

20 The PVB particles dispersed on the polymer surface were shown as the tiny holes in the following two Figure 1 and Figure 2. The tensile bar of Example E15 was extracted with methanol to remove surface PVB. SEM pictures were taken on two arbitrary surfaces after methanol extraction.

Examples E45 to 56 and Control Examples C7 to C10:
Sound Reduction Results for Laminated Glass Filled
Nylon 6 Round Disc.

5

The same process and procedures in above Examples
E12 to E14, & C2 were used for Examples E45 to E56 & C7
to C10 in Table 9 except round discs of 4 inch diameter
with 2mm thickness were molded in an injection molding
10 machine. Three types of PVB sheeting were used as the
interlayer to make laminated polymer discs:

1. Stiff PVB (less than 30 pph plasticizer)
2. Standard Butacite® B140C
- 15 3. Special PVB (43% plasticizer)

Those laminated polymer discs were used in a sound
reduction test unit as described in the following.

20 Sound Reduction Test

Sound intensity was measured from 12.5 to 10,000
Hz (Hertz) frequency at 12.5 Hz increments. Duplicate
results were recorded for the incoming sound (S), that
25 is the sound prior to attenuation by the laminate. The
sound intensity passing through a laminated polymer
plate (S*) was also recorded. The absorption intensity
or the sound power level loss (SPLL) in dB at each
frequency was calculated according to the following
30 equation:

$$SPLL = 20 * \text{LOG}_{10} (S^*/S_{\text{Avg}}),$$

wherein S_{Avg} is the average of two incoming sound
measurements.

35

The sum of all frequencies are recorded in Table 9
under the column heading "**Total Sound Reduction in dB.**"

For the above Sound Reduction Test, laminated
40 polymer discs were used for E45 to E56, while two non-
laminated discs were used for the control examples.
The difference in total sound reduction for a laminated
disc versus the control with two non-laminated discs
was also recorded in Table 9 under column heading "dB
45 Reduction versus Control." The sound power level loss
in dB was also plotted versus sound frequency as shown

in Figure 3 for C9, E51, E52, & E53. The sound reduction is much better for laminated polymer discs than the polymer without lamination, especially in the frequency ranges of 1000 to 2000 Hz, 4000 to 6500 Hz, and 8000 to 9500 Hz.

Table 9
Summary of Sound Intensity Reduction

Example #	Additive (13 Wt%)	Interlayer	Total Sound Reduction in dB	dB Reduction versus Control
C7	ECOCITE™ E	None	-40.63	Control 1^a
E45	ECOCITE™ E	1	-44.05	-3.41
E46	ECOCITE™ E	2	-45.80	-5.16
E47	ECOCITE™ E	3	-45.76	-5.13
C8	ECOCITE™ G	None	-42.50	Control 2^b
E48	ECOCITE™ G	1	-44.25	-1.75
E49	ECOCITE™ G	2	-46.27	-3.76
E50	ECOCITE™ G	3	-46.60	-4.09
C9	ECOCITE™ H	None	-41.85	Control 3^c
E51	ECOCITE™ H	1	-44.92	-3.06
E52	ECOCITE™ H	2	-45.79	-3.94
E53	ECOCITE™ H	3	-46.19	-4.34
C10	T-mix	None	-40.96	Control 4^d
E54	T-mix	1	-43.72	-2.76
E55	T-mix	2	-45.90	-4.94
E56	T-mix	3	-45.58	-4.62

^aControl sample for E45 to E47.

^bControl sample for E48 to E50.

^cControl sample for E51 to E53.

^dControl sample for E54 to E56.

Example 57 is a trilayer laminate of stiff (<30 pph plasticizer) PVB sandwiched between two sheets of Glass-filled Nylon 6 with added Ecocite™ H (13 wt%). C11 is two sheets of glass-filled Nylon 6 sheet with Ecocite™ H, not laminated.

Example 58 is similar to E57 except that the Nylon 6 is not glass-filled, and the Ecocite™ is included at a level of 30 wt%. C12 is similar to C11 except that Nylon 6 is not glass-filled.

A pin was projected at each target from a load cell in a tower at a predetermined velocity. The energy required to penetrate the target was measured and recorded in Table 10. The percent increase for each example of the invention was determined by subtracting the total impact of the comparative example from the total impact of the example, and dividing the

difference by the total impact of the comparative example, then multiplying by 100%.

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Table 10

Example	Max Load (J)	Impact (J)	Velocity (m/sec)	Total (J)	Increase (%)
C11	6.6	119.3	2.3	25.7	--
E57	11.7	118.8	2.2	39.9	55.1
C12	66.2	118.9	2.3	112.5	--
E58	91.4	118.6	2.3	125.2	11.3